

FORM PTO-1390  
(REV 10-2000)

U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE

ATTORNEY'S DOCKET NUMBER

TRANSMITTAL LETTER TO THE UNITED STATES  
DESIGNATED/ELECTED OFFICE (DO/EO/US)  
CONCERNING A FILING UNDER 35 U.S.C. 371

0730.0040C

U.S. APPLICATION NO. (If known, see 37 CFR 1.5)

09/936781

INTERNATIONAL APPLICATION NO.  
PCT/AU00/00208INTERNATIONAL FILING DATE  
March 17, 2000PRIORITY DATE CLAIMED  
March 19, 1999

TITLE OF INVENTION PROCESS FOR THE REMOVAL OF OXALATE AND/OR SULPHATE FROM BAYER LIQUORS

APPLICANT(S) FOR DO/EO/US Steven P. Rosenberg et al.

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. ☒ This is an express request to promptly begin national examination procedures (35 U.S.C. 371(f)).
4. ☒ The US has been elected by the expiration of 19 months from the priority date (PCT Article 31).
5. ☒ A copy of the International Application as filed (35 U.S.C. 371(c)(2))
  - a. ☒ is attached hereto (required only if not communicated by the International Bureau).
  - b. ☒ has been communicated by the International Bureau.
  - c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US).
6. ☐ An English language translation of the International Application as filed (35 U.S.C. 371(c)(2)).
7. ☐ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))
  - a. ☐ are attached hereto (required only if not communicated by the International Bureau).
  - b. ☐ have been communicated by the International Bureau.
  - c. ☐ have not been made; however, the time limit for making such amendments has NOT expired.
  - d. ☐ have not been made and will not be made.
8. ☐ An English language translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
9. ☒ An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).
10. ☐ An English language translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).

## \* Items 11 to 16 below concern document(s) or information included:

11. ☐ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
12. ☒ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
13. ☐ A **FIRST** preliminary amendment.  
☐ A **SECOND** or **SUBSEQUENT** preliminary amendment.
14. ☐ A substitute specification.
15. ☐ A change of power of attorney and/or address letter.
16. ☒ Other items or information:

Copy of International Preliminary Examination Report; Copy of Notification of Information Concerning Elected Offices Notified of Their Election; and a Postcard Receipt.

U.S. APPLICATION NO. (If known, see 37 CFR 1.5)

09/936781

INTERNATIONAL APPLICATION NO  
PCT/AU00/00208

ATTORNEY'S DOCKET NUMBER  
0730.0040C

17. ☒ The following fees are submitted:

**BASIC NATIONAL FEE ( 37 CFR 1.492 (a) (1) - (5) ) :**

Neither international preliminary examination fee (37 CFR 1.482)  
nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO  
and International Search Report not prepared by the EPO or JPO . . . . . **\$1000.00**

International preliminary examination fee (37 CFR 1.482) not paid to  
USPTO but International Search Report prepared by the EPO or JPO . . . . . **\$860.00**

International preliminary examination fee (37 CFR 1.482) not paid to USPTO but  
international search fee (37 CFR 1.445(a)(2)) paid to USPTO . . . . . **\$710.00**

International preliminary examination fee paid to USPTO (37 CFR 1.482)  
but all claims did not satisfy provisions of PCT Article 33(1)-(4) . . . . . **\$690.00**

International preliminary examination fee paid to USPTO (37 CFR 1.482)  
and all claims satisfied provisions of PCT Article 33(1)-(4) . . . . . **\$100.00**

**ENTER APPROPRIATE BASIC FEE AMOUNT =**

**CALCULATIONS** PTO USE ONLY

\$ 1000.00

Surcharge of **\$130.00** for furnishing the oath or declaration later than ☐ 20 ☐ 30  
months from the earliest claimed priority date (37 CFR 1.492(e)).

\$

CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE
Total claims	20 - 20 =	0	X <b>\$18.00</b>
Independent claims	2 - 3 =	0	X <b>\$80.00</b>
MULTIPLE DEPENDENT CLAIM(S) (if applicable)			+ <b>\$270.00</b>

\$ 0.00

\$ 0.00

\$ 0.00

**TOTAL OF ABOVE CALCULATIONS =**

\$ 1,000.00

☒ Applicant claims small entity status. See 37 CFR 1.27. The fees indicated above  
are reduced by 1/2.

\$

**SUBTOTAL =**

\$ 1,000.00

Processing fee of **\$130.00** for furnishing the English translation later than ☐ 20 ☐ 30  
months from the earliest claimed priority date (37 CFR 1.492(f)).

\$ 00.00

**TOTAL NATIONAL FEE =**

\$ 1,000.00

Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be  
accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). **\$40.00** per property

\$ 40.00

**TOTAL FEES ENCLOSED =**

\$ 1,040.00

Amount to be  
refunded:

\$

charged:

\$

a. ☒ A check in the amount of \$ 1,040.00 to cover the above fees is enclosed.

b. ☐ Please charge my Deposit Account No. \_\_\_\_\_ in the amount of \$ \_\_\_\_\_ to cover the above fees.  
A duplicate copy of this sheet is enclosed.

c. ☒ The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any  
overpayment to Deposit Account No. 05-0460. A duplicate copy of this sheet is enclosed.

**NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.**

SEND ALL CORRESPONDENCE TO:

Finnan, Patrick J.  
EPSTEIN, EDELL, SHAPIRO, FINNAN & LYTLE, LLC  
1901 Research Blvd. Suite 400  
Rockville, Maryland 20850 US



SIGNATURE:

Patrick J. Finnan

NAME

39,189

REGISTRATION NUMBER

PROCESS FOR THE REMOVAL OF OXALATE AND/OR  
SULPHATE FROM BAYER LIQUORS

FIELD OF THE INVENTION

- 5 The present invention relates to a process and apparatus for the removal and causticisation of sodium oxalate and/or sodium sulphate from a Bayer process liquor.

BACKGROUND TO THE INVENTION

- 10 In the Bayer process for the production of alumina, bauxite is digested in a caustic liquor, generally under conditions of elevated temperature and pressure. A variety of organic and inorganic impurities are invariably extracted at the same time, reacting with caustic soda to form their sodium salts. In addition, some of the organic compounds can undergo degradation, ultimately producing sodium carbonate and the sodium salts of a range of simple carboxylic acids. The formation of these impurities represents a major loss of
- 15 caustic from the refinery's liquor streams. This caustic must either be replaced, or recovered in some way from the impurities.

- The recovery of caustic from sodium carbonate is a commonplace activity in most alumina refineries. The causticisation of sodium carbonate is generally effected by the addition of
- 20 lime, which reacts with the sodium carbonate to form calcium carbonate, thereby liberating sodium hydroxide. An improved version of this process is described in our co-pending International Application No. PCT/AU99/00757, filed on the 25<sup>th</sup> of September 1999 and entitled "Improved Bayer Causticisation". The contents of PCT/AU99/00757 are incorporated herein by reference.

25

- Of the other impurities, sodium oxalate and sodium sulphate are among the most significant. The presence of sodium oxalate in Bayer process streams is problematical owing to its very limited solubility. This creates a number of well-known problems within the alumina refinery. Sodium sulphate is much more soluble, and can accumulate to very
- 30 high concentrations. This causes a different set of problems, particularly with respect to the refinery's productivity. The problems associated with this impurity in Bayer process liquors, and a process for its separation, have been described in Australian patent No. 673306.

Many prior art processes have been described for the removal of sodium oxalate and sodium sulphate from Bayer liquors. Some of these processes remove both impurities concurrently. In most cases, these processes advocate that the impurity is discarded after removal from the liquor stream. However, a small number of the above processes also provide a means for the recovery of soda from sodium oxalate. None describe a practical method for the recovery of soda from sodium sulphate, requiring that it be discarded. However, disposal of sodium sulphate is not straightforward.

Environmental considerations preclude disposal of sodium sulphate into natural water systems, and since it is highly soluble, it must be disposed in a suitably lined or otherwise isolated sanitary landfill if it is not to enter groundwater systems. In the alumina refinery, disposal of sodium sulphate to the red mud residue disposal areas results in the eventual return of most of the sodium sulphate to the process liquor stream with the recovered lake water.

Whilst it is preferable to utilise the sodium sulphate in some way, for example by conversion into useful products, options for this are extremely limited. Electrolytic cells are commercially available which convert sodium sulphate into sodium hydroxide and either sodium bisulphate or sulphuric acid. However, these are generally restricted to reasonably pure solutions in which scales are unlikely to form, because the membranes used in the cells are sensitive to fouling. Other processes have been investigated including reductive processes such as the Leblanc process, and the Peniakoff process for production of gibbsite from bauxite. These latter processes are not currently practised, as they are inefficient, costly and produce environmentally unacceptable by-products.

Thus, there is a significant need for an economic process for the processing of sodium sulphate into more useful products, and/or for the immobilisation of the sulphate anion in an environmentally acceptable, insoluble material.

Most alumina refineries practice some form of oxalate removal process. In general, these processes are based on variations of the following two procedures:

1. Sodium oxalate is permitted to coprecipitate with gibbsite in the refinery's gibbsite

precipitation circuit. The co-crystallised oxalate reports to the refinery's gibbsite seed preparation facility, where it is removed by washing with water or dilute liquor. The oxalate-rich washings are then further treated to remove oxalate either by seeding and evaporation to recrystallise sodium oxalate or, by reaction with lime, as calcium oxalate.

5

2. Oxalate co-crystallisation is avoided by crystallising and removing sodium oxalate in a side-stream of one of the refinery's main process streams (usually a spent liquor stream).

The side stream is evaporated to increase the supersaturation of the sodium oxalate and directed to a series of oxalate crystallisers where it is seeded with recycled sodium oxalate

10 crystals. After solid/liquid separation, the clarified and now oxalate-depleted liquor is returned to the process. A portion of the solid sodium oxalate is recycled to act as seed, while the remainder is either discarded or processed to recover soda. An example of this process is outlined in US 3,899,571.

15 Most processes for the recovery of the soda values from sodium oxalate are based on reactions with lime. In some processes, the separated sodium oxalate cake is first burnt in a kiln to produce sodium carbonate, which is subsequently causticised by reaction with lime. This process is costly to operate, and the conversion to sodium carbonate is not always complete.

20

In other processes, a solution rich in sodium oxalate, such as the washings from the seed circuit of a refinery that practices coprecipitation of oxalate, is directly reacted with lime to form calcium oxalate. However, whilst very low oxalate concentrations can be achieved in the treated stream in this way, the efficiency of lime utilisation is very poor, due to the  
25 formation of calcium aluminates such as tricalcium aluminate (TCA), unless the stream is very low in caustic and sodium aluminate. Consequently, this process can only be applied to dilute liquors.

### SUMMARY OF THE INVENTION

30 The present invention was developed with a view to providing a means for the direct removal of sodium sulphate or sodium oxalate, or combinations of both, in Bayer process liquors with the production of sodium hydroxide. The unwanted anion is isolated in an insoluble solid material that can be disposed of in a conventional sanitary landfill, thus

preventing the return of the unwanted anions to the refinery via the refinery's lake system.

Throughout this specification, we have used conventional North American terminology for the description of Bayer solution compositions. Thus, 'C' refers to the caustic concentration of the liquor, this being the sum of the sodium aluminate and sodium hydroxide content of the liquor expressed as equivalent g/L of sodium carbonate. 'S' refers to the sum of C and the true concentration of sodium carbonate. Thus, S-C gives the actual concentration of  $\text{Na}_2\text{CO}_3$  in the liquor, in g/L. 'A' refers to the concentration of sodium aluminate in the liquor, expressed as equivalent g/L of  $\text{Al}_2\text{O}_3$ .

10

Sodium oxalate concentration is expressed as g/L of  $\text{Na}_2\text{C}_2\text{O}_4$ . Sodium sulphate concentration is expressed as g/L of  $\text{Na}_2\text{SO}_4$ . 'TS' refers to the sum of all sodium salts in solution, expressed as the equivalent concentration in g/L of sodium carbonate.

15 Lime refers either to calcium oxide, or more preferably, calcium hydroxide. Lime efficiency is defined as the percentage ratio of the number of moles of sodium hydroxide produced to the number of moles of lime consumed, divided by two.

The term Hydrocalumite is used to refer to any layered double hydroxide compound formed between calcium and aluminium, within which charge balancing anions are intercalated. Typically, these compounds will be of the form  $[\text{Ca}_2\text{Al}(\text{OH})_6]_2 \cdot \text{X} \cdot n\text{H}_2\text{O}$ , where 'X' represents a charge-balancing anion or anions.

Throughout this specification the term "comprising" is used inclusively, in the sense that there may be other features and/or steps included in the invention not expressly defined or comprehended in the features or steps subsequently defined or described. What such other features and/or steps may include will be apparent from the specification read as a whole.

According to one aspect of the present invention there is provided a process for the removal and causticisation of sodium oxalate and/or sodium sulphate from a Bayer process liquor containing sodium carbonate and one or both of sodium oxalate and sodium sulphate in an alumina refinery, the process comprising the steps of:

removing aluminate ions from the Bayer liquor through the formation of a carbonate-bearing hydrocalumite and/or sulphate-bearing hydrocalumite; and,

5 treating the liquor with sufficient lime to remove and causticise any residual carbonate ions and some or all of the oxalate ions present whereby any reacted lime solids thus formed can be separated and safely disposed of.

Preferably the process comprises a further step, prior to said step of removing aluminate ions, in which the liquor is enriched with sulphate and/or oxalate such that any aluminate  
10 and/or carbonate ions entering with the sulphate and/or oxalate are also removed.

Preferably the process comprises a further step, following said step of removing aluminate ions, of separating the carbonate-bearing hydrocalumite and/or sulphate-bearing hydrocalumite from the Bayer liquor to form a clarified liquor.

15 In another embodiment the process further comprises a pre-causticisation step in which the Bayer liquor is first causticised to reduce the concentration of carbonate ions, prior to said step of removing aluminate ions. Typically said pre-causticisation step includes heating the liquor, adding sufficient lime to react with the carbonate ions to produce calcium  
20 carbonate and separating the calcium carbonate from the liquor. Optionally the heated liquor is enriched with sulphate and/or oxalate prior to causticisation to ensure that any carbonate ions entering with the sulphate and/or oxalate are also causticised.

According to another aspect of the present invention there is provided an apparatus for the  
25 removal and causticisation of sodium oxalate and/or sodium sulphate from a Bayer process liquor containing sodium carbonate and one or both of sodium oxalate and sodium sulphate in an alumina refinery, the apparatus comprising:

30 means for removing aluminate ions from the Bayer liquor through the formation of a carbonate-bearing hydrocalumite and/or sulphate-bearing hydrocalumite; and,

means for treating the liquor with sufficient lime to remove and causticise any residual carbonate ions and some or all of the oxalate ions present whereby any

reacted lime solids thus formed can be separated and safely disposed of.

In one embodiment said means for removing aluminate ions comprises a first reaction vessel to which sufficient lime is added to react with all of the aluminate ions in the liquor.

5 Preferably the apparatus of this embodiment further comprises a means for separating the carbonate-bearing hydrocalumite and/or sulphate-bearing hydrocalumite from the liquor to form a clarified liquor. Preferably the means for treating the liquor comprise a second reaction vessel to which sufficient lime is added to react with the sodium oxalate in the clarified liquor to form calcium oxalate and with any remaining carbonate ions to form  
10 sodium carbonate.

In another embodiment said means for removing aluminate ions and said means for treating the liquor are comprised in a single reaction vessel to which sufficient lime is added to react with the aluminate ions to form said carbonate-bearing and/or sulphate-  
15 bearing hydrocalumite, together with sufficient additional lime to react with the sodium oxalate to form calcium oxalate.

The inventors' theories on the chemical reactions in the process and apparatus of the present invention are merely examples of possible reactions thought to be taking place and  
20 are not intended to be limiting in any way.

### **BRIEF DESCRIPTION OF DRAWINGS**

In order to facilitate a better understanding of the nature of the invention preferred embodiments of the process and apparatus for the removal and causticisation of sodium  
25 oxalate and/or sodium sulphate will now be described in detail, by way of example only, with reference to the accompanying drawings in which:

Figure 1 is a simplified process flow diagram for the causticisation of sodium oxalate in accordance with one embodiment of the present invention; and,  
30

Figure 2 is a simplified process flow diagram for the causticisation and removal of sodium sulphate and sodium oxalate in accordance with another embodiment of the present invention.

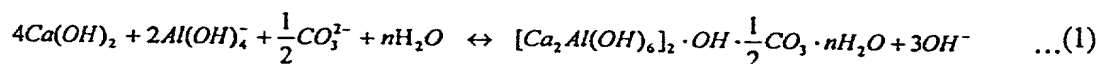


### DETAILED DESCRIPTION OF PREFERRED EMBODIMENTS

This invention is based upon the following two key findings:

1. The inventors have found that lime will not react with sodium oxalate to form calcium oxalate and sodium hydroxide in Bayer process liquors unless the concentration of sodium aluminate is close to zero.
2. It is known that hydrocalumite-type structures are formed by the reaction of lime with sodium aluminate solutions such as Bayer liquors, and that these structures will incorporate anions into the interlayer regions. The order of preference by which anions are incorporated is critical in the utilisation of this property. The inventors have found that the order of preference of anion incorporation in Bayer liquors is carbonate > sulphate > oxalate. This contrasts with published literature which indicates that the order is carbonate > oxalate > sulphate. ["Layered Double Hydroxides for Treatment of Bayer Process Lake Water" AJ Perrotta, FS Williams and LC Stonehouse, Light Metals (1997), 37-48].

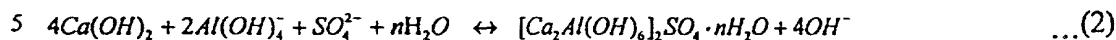
The above two findings explain why the conventional approaches to causticising sodium oxalate in Bayer process solutions are so inefficient with respect to lime use, and must be restricted to low S liquors. The addition of lime to these solutions results in the formation of hydrocalumite (Hc). Since carbonate is usually present in these liquors, the Hc formed is primarily the carbonate form:



If sufficient lime has been added, this reaction will proceed until virtually all of the aluminate ion in solution has been removed, other than a small equilibrium concentration. Note that this is a very inefficient causticising reaction, in that eight moles of  $\text{Ca}(\text{OH})_2$  are required to causticise one mole of  $\text{Na}_2\text{CO}_3$ .

Rarely, insufficient carbonate may be present in the liquor to counterbalance all of the Hc.

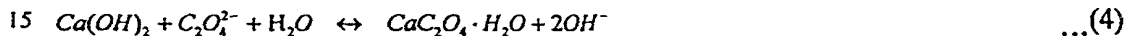
In this case, if sulphate and/or oxalate are also present, sulphate is preferentially incorporated into the inter-layer regions of the hydrocalumite structure, as indicated by the following equation:



If insufficient sulphate is present to counterbalance all of the Hc formed, a small amount of oxalate can be incorporated into the Hc structure, as follows:



Once the aluminate has been reduced to close to its equilibrium concentration in contact with Hc, the addition of further lime will result in the following reaction:



Reaction (4) is preferred over reaction (3), since only one mole of calcium hydroxide is required to causticise each mole of oxalate, and no alumina is consumed.

20 The reasons for the poor lime efficiency of conventional sodium oxalate causticisation processes should be apparent from the above explanation to those skilled in the arts of the Bayer process. Most of the lime is consumed forming Hc according to equation (1) above. When the dissolved alumina has been consumed, any additional lime will react with the oxalate and residual carbonate to form calcium oxalate and calcium carbonate  
25 respectively. Consequently, since most Bayer liquors contain substantial dissolved alumina, and the amount is usually proportional to the S concentration of the liquor, it is clear why the process is usually restricted to solutions of low S concentration.

From the preceding discussion, it is apparent that to efficiently causticise sodium oxalate  
30 solutions, it is first necessary to remove the aluminate ion from solution, preferably with recovery of the aluminate ion in some later step. In this invention, this removal is effected

by reacting the aluminate with lime in such a manner that productive use is made of the hydrocalumite so formed. In its preferred form the process consists of the following basic steps:

- 5 1. Removal of carbonate ions from a Bayer liquor containing sodium carbonate and one or both of sodium oxalate and sodium sulphate, either through the formation of calcium carbonate using any suitable carbonate causticisation process known to those skilled in the arts of the Bayer process, or through the formation of carbonate-bearing hydrocalumite according to equation (1) above.
- 10 2. Separation and disposal of any calcium carbonate formed in Step (1) above, or separation and recovery of any carbonate-bearing hydrocalumite formed in Step (1). The recovered Hc may then be used for further causticisation, using the process described in co-pending International Application No. PCT/AU99/00757, thereby  
15 recovering the alumina consumed in Step (1).
3. Treatment of the clarified liquor produced in Step (2) with sufficient lime to remove any dissolved alumina remaining after Step (1), plus enough additional lime to react any remaining sodium carbonate and the sodium oxalate. This results in the  
20 following sequence of reactions:
  - the removal and causticisation of some or all of the residual carbonate as carbonate-bearing Hc according to equation (1);
  - the removal and causticisation of some or all of the sodium sulphate as sulphate-bearing Hc according to equation (2), and;
  - 25 the removal and causticisation of sodium oxalate as predominantly calcium oxalate monohydrate according to equation (4). Some calcium carbonate may also form.
4. Separation and disposal of the reacted lime solids formed in Step (3), and return of  
30 the clarified caustic solution to a suitable location within the Refinery.

The process of the invention for the causticisation and removal of sodium oxalate and/or sodium sulphate is further described and illustrated in the following two examples. These examples are illustrative of a variety of possible implementations and are not to be construed as limiting the invention in any way.

5

### EXAMPLE 1

In this example, oxalate is causticised to sodium hydroxide with high lime efficiency and with little loss of alumina. The process may be operated in either batch or continuous mode with suitable selection of equipment.

10

Referring to the simplified process flow diagram of Figure 1, a Bayer process liquor of S concentration of between 0 and 250 g/L, preferably less than 150 g/L is directed into a reaction vessel 10 and maintained at a temperature of between 20 and 90°C, preferably between 50 and 70°C. The type of reactor is not critical, for example a CSTR may be  
15 used, provided that sufficient agitation is applied to ensure that all of the solids are adequately suspended. If the solution is not already enriched with oxalate, sodium oxalate in either solid form or as an aqueous solution may optionally be added, provided that the solubility of sodium oxalate is not exceeded after mixing with the liquor to be treated. A suitable stream for treatment would be the filtrate from the gibbsite seed washing facility  
20 in a refinery practising oxalate co-precipitation. Alternatively, a suitable stream could be prepared by dissolving oxalate cake in a dilute liquor such as the filtrate from the product washing filters.

Sufficient lime (preferably slaked lime) is added to the reaction vessel to react with  
25 substantially all of the dissolved alumina in the solution, forming a carbonate-bearing hydrocalumite. The lime requirement can be calculated using equation (1) above. The residence time in this reactor is not critical. The reaction is generally found to be complete in less than five minutes, but residence times of up to 2 hours have little or no adverse effect. The preferred residence time is 30 minutes. Excessive residence times may result in  
30 the undesirable formation of TCA, especially at high temperatures, causing a loss of efficiency.

The hydrocalumite solids and liquor are then separated using any suitable solid/liquid separation device 12 (preferably a pressure filter). The solids may then be used to causticise another liquor stream within the refinery, using the process revealed in co-  
5 pending International Application No. PCT/AU99/00757.

The clarified liquor is then directed to a second reaction vessel 14 and sufficient lime (preferably slaked lime) is added to react with the sodium oxalate to form calcium oxalate, and with any remaining sodium carbonate to form calcium carbonate. The amount of lime required may be calculated using equation (4) above, together with the following equation:  
10



This reaction should be conducted between 20 and 140°C, preferably between 50 and 80°C, with a reaction time of between 15 mins and 4 hours, preferably 60 minutes.  
15

The resultant slurry is then forwarded to any suitable solid/liquid separation device 16, preferably a pressure filter. The solids may be discharged to the alumina refinery's red mud disposal area, or after washing and drying, calcined for re-use.

20

### Laboratory Test Results

#### STAGE 1 (a) - Removal of Aluminate Ions:

A calcium hydroxide slurry was prepared by slaking 31.3g of freshly calcined LR grade CaO in 350 mL of deionised water. 860 mL of simulated seedwash filtrate liquor was  
25 transferred to a 2 litre stainless steel Parr autoclave and heated to 60°C. The temperature was maintained thermostatically. Agitation was applied using a pitched blade turbine impeller rotating at 200 rpm. When the temperature of the system had equilibrated, the slaked lime slurry was added quantitatively.

30 Samples of the liquors were collected from the reactor and filtered using 0.45µm Acrodisc filters, at the commencement of the test and after 10 minutes of reaction.

After 10 minutes of reaction, the agitator was stopped, the slurry removed from the autoclave and filtered under vacuum through a Whatmans #1 filter. This step was performed rapidly to avoid carbonation of the liquor by reaction with carbon dioxide in the  
5 air.

#### STAGE 1(b) - Use of Hydrocalumite for Causticisation:

The hydrocalumite solids (341g) collected by the filter were washed and air-dried. A sample of liquor was collected from the refinery mud washing circuit and filtered through  
10 a Whatman's #1 filter paper. 1000 mL of this liquor was placed in the autoclave and heated to 100°C. The temperature was maintained thermostatically. Agitation was applied using a pitched blade turbine impeller rotating at 200 rpm. When the temperature of the system had equilibrated, 80.4 g of the hydrocalumite solids was added quantitatively. The reaction was allowed to proceed for two hours with periodic sampling throughout.

#### STAGE 2 - Oxalate Removal

A portion of the filtrate (670 mL) from Stage 1(a) was returned to the autoclave and equilibrated at a temperature of 60°C. The agitator was restarted and operated at 200 rpm. Calcium hydroxide slurry, prepared by slaking 6.2g of freshly calcined LR grade CaO in  
20 70 mL of deionised water, was then quantitatively added to the reactor.

Samples of the liquors were collected from the reactor and filtered using 0.45 $\mu$ m Acrodisc filters, just prior to addition of the lime slurry and after 30 minutes of reaction. At 30 minutes, the agitator was stopped, the slurry removed from the autoclave and filtered  
25 under vacuum through a Whatmans #1 filter paper. As before, this step was performed rapidly to avoid carbonation of the liquor by carbon dioxide in air.

#### Comparison With Prior Art

To compare the efficiency of the proposed process with the prior art processes, a sample of  
30 the preceding liquor was reacted in a single stage process with slaked lime.

A sample of the simulated seed-wash filtrate liquor above (900mL) was transferred to a 2 litre stainless steel Parr autoclave and heated to 60°C. Calcium hydroxide slurry was prepared by slaking 52.9g of freshly calcined LR grade CaO in 550 mL of deionised water. The temperature was maintained thermostatically. Agitation was applied using a pitched blade turbine impeller rotating at 200 rpm. When the temperature of the system had equilibrated, the slaked lime slurry was added quantitatively.

Samples of the liquors were collected from the reactor and filtered using 0.45 $\mu$ m Acrodisc filters, at the commencement of the test and after 30 minutes of reaction.

- 10 After 30 minutes of reaction, the agitator was stopped, the slurry removed from the autoclave and filtered under vacuum through a Whatmans #1 filter. This step was performed rapidly to avoid carbonation of the liquor by reaction with carbon dioxide in the air.

## 15 Results:

Table 1: Liquor Analyses for Stage 1A of process.

Sample	Liquor Volume	A (g/L)	C (g/L)	S (g/L)	A/C	C/S	Na <sub>2</sub> CO <sub>3</sub> (g/L)	NaCl (g/L)	Na <sub>2</sub> SO <sub>4</sub> (g/L)	Na <sub>2</sub> C <sub>2</sub> O <sub>4</sub> (g/L)	TS (g/L)
Seedwash filtrate (t=0)	860 mL	19.8	55.6	66.4	0.356	0.837	10.8	5.4	15.1	8.8	89.8
t = 10 mins	1197 mL	2.8	51.6	52.3	0.054	0.987	0.7	3.4	8.2	5.3	64.5

Analysis of the solids by XRD indicated predominantly carbonate-bearing

- 20 hydrocalumite phases (major), some sulphate-bearing hydrocalumite (minor), a small amount of calcium oxalate (minor) and calcium carbonate (trace).

Table 2: Liquor Analyses for Stage 1B of process.

Sample	Liquor Volume	A (g/L)	C (g/L)	S (g/L)	A/C	C/S	Na <sub>2</sub> CO <sub>3</sub> (g/L)	NaCl (g/L)	Na <sub>2</sub> SO <sub>4</sub> (g/L)	Na <sub>2</sub> C <sub>2</sub> O <sub>4</sub> (g/L)	TS (g/L)
Mud Washer overflow (t=0)	1000 mL	66.5	103.9	126.7	0.640	0.820	22.8	9.0	24.2	1.8	174.3
t = 120 mins	1024 mL	68.0	109.9	123.8	0.619	0.888	13.9	9.0	24.3	2.5	170.2

Analysis of the solids by XRD indicated predominantly calcium carbonate (major phase) with some carbonate-bearing hydrocalumite (trace).

As can be seen from the results in the above two tables, sodium carbonate has been efficiently removed and caustic generated. Small amounts of sodium sulphate and sodium oxalate have also been causticised.

Table 3: Liquor Analyses for 2<sup>nd</sup> stage of process

Sample	Liquor Volume	A (g/L)	C (g/L)	S (g/L)	A/C	C/S	Na <sub>2</sub> CO <sub>3</sub> (g/L)	NaCl (g/L)	Na <sub>2</sub> SO <sub>4</sub> (g/L)	Na <sub>2</sub> C <sub>2</sub> O <sub>4</sub> (g/L)	TS (g/L)
Seedwash filtrate (t=0)	670 mL	3.1	51.6	52.4	0.060	0.985	0.8	3.4	8.3	5.2	65.6
t = 30 mins	753 mL	0.3	51.3	51.8	0.006	0.990	0.5	2.6	4.5	1.5	58.4

Analysis of the solids by XRD indicated sulphate-bearing hydrocalumite (major), calcium oxalate (major) and unreacted lime (trace).

It can be seen from the above results that the concentration of oxalate and

sulphate are greatly reduced. Small amounts of carbonate and other impurities have also been removed.

#### Prior Art Process

Table 4: Liquor Analyses for Prior Art simulation.

Sample	Liquor Volume	A (g/L)	C (g/L)	S (g/L)	A/C	C/S	Na <sub>2</sub> CO <sub>3</sub> (g/L)	NaCl (g/L)	Na <sub>2</sub> SO <sub>4</sub> (g/L)	Na <sub>2</sub> C <sub>2</sub> O <sub>4</sub> (g/L)	TS (g/L)
Seedwash filtrate (t=0)	900 mL	20.0	55.2	66.1	0.362	0.835	10.9	4.9	14.9	10.10	89.1
t = 30 mins	1387 mL	0.2	51.8	52.4	0.004	0.989	0.6	2.8	5.1	0.9	57.8

Analysis of the solids by XRD indicated multiple hydrocalumite phases (due to

the presence of several different counterbalancing ions), calcium oxalate (major)



and some unreacted lime (trace).

### Comparison of Example 1 with Prior Art

After due allowance for volumetric changes due to the input of water with slaked  
5 lime, evaporative losses and changes in liquor composition, the following  
performance results were obtained for Stages 1 and 2 of Example 1, and for the  
simulation of the Prior Art process.

Table 5: Comparison of Causticisation and Removal  
for Example 1 with Prior Art Process.

	Stage 1 t/t CaO	Stage 2 t/t CaO	Prior Art t/t CaO
Alumina loss (as $\text{Al}_2\text{O}_3$ )	0	-0.298	-0.335
Sodium carbonate converted	-1.491	-0.026	-0.170
Sodium sulphate converted	-0.003	-0.351	-0.120
Sodium oxalate converted	0	-0.379	-0.148
Sodium chloride converted	0	-0.052	-0.009
Sodium hydroxide produced	1.269	0.492	0.317
Lime efficiency	88.9%	34.4%	22.2%

The average lime efficiency for Example 1 over both stages of the process was  
77.9%.

It can be seen from the above results that the proposed process is considerably more  
efficient in terms of its lime utilisation, and that far greater removal of both oxalate and  
sulphate is effected per tonne of lime. Furthermore, it should be noted that the mass of

alumina lost per tonne of sodium oxalate removed is reduced by a factor of three.

### EXAMPLE 2

In this example, sodium carbonate, sodium sulphate and sodium oxalate may all be  
5 causticised and removed from solution. In essence, the process sacrifices dissolved  
alumina for the recovery of the soda values of the sodium sulphate. Consequently, loss  
of alumina is of the same magnitude as would have occurred in the prior art oxalate  
causticisation processes, but with far greater recovery of caustic soda. While recovery  
of the alumina is feasible, it cannot be done without attendant release of the sulphate,  
10 and is hence not discussed here.

The principles of this variant of the process are best described by reference to the  
simplified flow diagram shown in Figure 2. The process may be operated in either  
batch or continuous mode with suitable selection of equipment.

15

A Bayer process liquor of S concentration of between 0 and 250 g/L, preferably less  
than 150 g/L is first causticised to reduce the carbonate concentration, using any  
suitable procedure known to those practised in the arts of the Bayer process.  
Preferably, the process disclosed in co-pending International Application No.  
20 PCT/AU99/00757 is used for this function, as this will ensure highest efficiency. In the  
example given here, the liquor is heated to close to the atmospheric boiling point of the  
liquor (approximately 100°C) by passing it through a heat exchanger 18 and directed  
into a reaction vessel 20. The type of reactor is not critical. For example, a CSTR may  
be used, provided that sufficient agitation is applied to ensure that all of the solids are  
25 adequately suspended.

If the solution is not already enriched with sulphate and/or oxalate, these may be added  
into this tank. Sodium oxalate may be added in either solid form or as an aqueous  
solution, provided that the solubility of sodium oxalate is not exceeded. The sodium  
30 sulphate can be prepared using any of the techniques described earlier and added either  
in solid form or as an aqueous solution, once again ensuring that the solubility of

anhydrous sodium sulphate is not exceeded. For example, a liquor stream fortified with sulphate suitable for treatment by this process is produced using the process described in Australian patent No. 673306.

- 5 While it is feasible to add these impurities into the second tank, or other suitable location, it is preferable if this is done in the first tank, or in the liquor stream prior to entering the first tank. This is to ensure that any sodium carbonate entering with the oxalate or sulphate is causticised, otherwise efficiency may be degraded.
- 10 Sufficient lime (preferably slaked lime) is added to the reaction vessel 20 to react with the sodium carbonate: the amount will depend upon the liquor composition and the causticisation technique employed. However, care should be taken to avoid overcharging of lime, as this will reduce the lime efficiency of the process. In the example given here, a residence time of approximately 1 hour was sufficient to ensure
- 15 optimum causticisation.

The slurry is then discharged to a solid/liquid separation device 22, which may be of any suitable design (preferably a pressure filter). The solids, which will consist primarily of calcium carbonate, may be discarded. Alternatively the solids may be

20 further washed to recover soda, and then calcined for re-use.

The clarified liquor is cooled to between 30 and 90°C, preferably between 50°C and 70°C in a heat exchanger 24 and directed to a second reaction vessel 26. The heat exchanger 24 may be located before solid liquid separation device 22 to make the

25 operating conditions within the solid/liquid separation device less aggressive. This will allow for a greater selection of solid/liquid separation devices. Once the liquid is separated and cooled, sufficient lime (preferably slaked lime) is added to react with the alumina, forming hydrocalumite, together with sufficient additional lime to react with the sodium oxalate to form calcium oxalate. The Hc thus formed will consist of a

30 mixture of both carbonate and sulphate-bearing species, depending upon the amount of residual carbonate remaining in the liquor. The carbonate-bearing species will form

preferentially, according to equation (1), followed by the sulphate-bearing species according to equation (2).

Thus, the ability of this process to causticise sodium sulphate will depend strongly on the efficiency of the pre-causticisation step in Tank 20, as well as the dissolved alumina content of the liquor, assuming that sodium sulphate is present in excess. Sodium oxalate will be causticised according to equation (4). The lime charge required can therefore be calculated on the basis of the liquor composition and the above three equations. To ensure efficient removal of the sodium oxalate, a slight excess of lime above the calculated amount (approximately 10%) is advisable.

The residence time required in the reactor 26 is between 30 minutes and 4 hours, preferably approximately 2 hours. Shorter residence times may be used if oxalate removal is not of paramount importance – in this case, the lime charge may be decreased accordingly.

The discharge from the tank 26 is pumped to a solid/liquid separation unit 28 such as a filter, centrifuge, or gravity separation device. The solids may then be disposed directly, or further washed and filtered before disposal. The washings can be returned to the second reactor, or mixed with the clarified liquor.

The clarified liquor, which will consist primarily of sodium hydroxide, may be returned to a suitable location within the refinery.

## 25 Laboratory Test Results

### STAGE 1:

A calcium hydroxide slurry was prepared by slaking 5.65g of freshly calcined LR grade CaO in 60 ml of deionised water. Refinery seed-wash filtrate liquor was collected and filtered through a Whatmans #1 filter paper. 1000mL of this filtered liquor was transferred to a 2 litre stainless steel Parr autoclave and heated to 100°C. The temperature was maintained thermostatically. Agitation was applied using a pitched

blade turbine impeller rotating at 200 rpm. When the temperature of the system had equilibrated, the slaked lime slurry was added quantitatively.

Samples of the liquor were collected from the reactor and filtered using 0.45 $\mu$ m

5 Acrodisc filters, at the commencement of the test and after 60 minutes of reaction.

After 60 minutes of reaction, the agitator was stopped, the slurry removed from the autoclave and filtered under vacuum through a Whatmans #1 filter. This step was performed rapidly to avoid carbonation of the liquor by reaction with carbon dioxide in  
10 the air. The results of analysis of the liquor following Stage 1 are given below in Table 6.

Table 6: Liquor Analyses for 1<sup>st</sup> stage of process.

Sample	Liquor Volume	A (g/L)	C (g/L)	S (g/L)	A/C	C/S	Na <sub>2</sub> CO <sub>3</sub> (g/L)	NaCl (g/L)	Na <sub>2</sub> SO <sub>4</sub> (g/L)	Na <sub>2</sub> C <sub>2</sub> O <sub>4</sub> (g/L)	TS (g/L)
Seedwash filtrate (t=0)	1000	20.0	55.2	66.1	0.362	0.835	10.9	5.1	14.9	10.1	89.1
t=60 mins	1053	18.8	61.2	63.7	0.307	0.961	2.5	4.9	14.1	9.4	84.6

15

Analysis of the solids by XRD indicated predominantly calcium carbonate (major phase) with some carbonate-bearing hydrocalumite (minor phase).

#### STAGE 2:

20 The filtrate (860 mL) was returned to the autoclave and equilibrated at a temperature of 60°C. The agitator was restarted and operated at 200 rpm. A calcium hydroxide slurry prepared by slaking 44.2g of freshly calcined LR grade CaO in 450 mL of deionised water was then quantitatively added to the reactor.

25 Samples of the liquors were collected from the reactor and filtered using 0.45 $\mu$ m Acrodisc filters, just prior to addition of the lime slurry and after 2 hours of reaction. At two hours, the agitator was stopped, the slurry removed from the autoclave and filtered under vacuum

through a Whatmans #1 filter paper. As before, this step was performed rapidly to avoid carbonation of the liquor by carbon dioxide in air. The results of analysis of the liquor following Stage 2 of the process are given in Table 7 below.

5

Table 7: Liquor Analyses for 2<sup>nd</sup> stage of process

Sample	Liquor Volume	A (g/L)	C (g/L)	S (g/L)	A/C	C/S	Na <sub>2</sub> CO <sub>3</sub> (g/L)	NaCl (g/L)	Na <sub>2</sub> SO <sub>4</sub> (g/L)	Na <sub>2</sub> C <sub>2</sub> O <sub>4</sub> (g/L)	TS (g/L)
Seedwash filtrate (t=0)	860	19.5	63.5	65.9	0.306	0.933	2.4	4.9	15.0	10.0	89.0
t=60 mins	1293	0.8	56.1	55.6	0.012	0.993	0.5	2.6	0.8	1.4	59.2

Analysis of the solids by XRD indicated multiple hydrocalumite phases (due to the presence of several different counterbalancing anions) as the major component, calcium oxalate (major phase) and unreacted lime (trace).

10

It can be seen from the above results that the concentrations of carbonate, oxalate and sulphate are greatly reduced, and that the C/S ratio has increased. Some reduction in sodium chloride concentration has also occurred, although the reduction is minor. Similarly, a mass balance over soda indicates that small quantities of impurities other than

15 those analysed here have also been causticised and removed.

After due allowance for volumetric changes due to the input of water with slaked lime, evaporative losses and changes in liquor composition, the following performance results were calculated per tonne of CaO consumed:

20

25

Table 8: Causticisation and Removal Performance for Example 2 compared with Prior Art Process

	Example 2 t/t CaO	Prior Art t/t CaO
Alumina loss (as $\text{Al}_2\text{O}_3$ )	-0.326	-0.335
Sodium carbonate converted	-0.167	-0.170
Sodium sulphate converted	-0.244	-0.120
Sodium oxalate converted	-0.142	-0.148
Sodium chloride converted	-0.016	-0.009
Sodium hydroxide produced	0.394	0.317
Total lime efficiency	27.6%	22.2%

5

From the above description of several preferred embodiments and illustrative examples, it will be apparent that the process and apparatus for removal and causticisation of sodium oxalate and/or sodium sulphate has a number of advantages, including the following:

- 10 (i) it provides an effective process for the removal of sodium sulphate;
- (ii) for the first time it provides a practical method for the recovery of soda from sodium sulphate;
- (iii) the efficiency of lime utilisation can be dramatically increased from about 20% to 80% (if sulphate removal is not the objective);
- 15 (iv) the oxalate concentration of the processed liquor is substantially lower than can usually be achieved in processes based on sodium oxalate crystallisation;
- (v) the efficiency of lime utilisation is greater than prior art processes based on reactions of lime with oxalate-rich Bayer liquors;
- (vii) unlike oxalate removal processes based on sodium oxalate crystallisation, the
- 20 process is not appreciably affected by the presence of organic poisons. This obviates the need for special organic poison removal processes, and contributes to consistent oxalate removal;

- (viii) the process does not require the recycling of seed crystals and the associated equipment to achieve this;
- (ix) the process does not require the use of strong liquors, raw caustic solutions or evaporation to supersaturate sodium oxalate. This simplifies oxalate removal and contributes to improved consistency of oxalate removal;
- (x) unlike many prior art oxalate removal processes based on sodium oxalate crystallisation, the precipitated solids are consistent in their filtration and deliquoring characteristics, despite quite wide variations in solution composition. Liquor throughput and residual cake moisture can both be optimised; and,
- (xi) the process provides supplementary causticisation capacity, raising the C/S of the refinery's liquors.

Numerous variations and modifications to the process and apparatus will suggest themselves to persons skilled in the Bayer process arts in Alumina refineries, in addition to those already described, without departing from the basic inventive concepts. All such variations and modifications are to be considered within the scope of the present invention, the nature of which is to be determined from the foregoing description and the appended claims.

20

25

30



THE CLAIMS DEFINING THE INVENTION

1. A process for the removal and causticisation of sodium oxalate and/or sodium sulphate from a Bayer process liquor containing sodium carbonate and one or both of sodium oxalate and sodium sulphate in an alumina refinery, the process comprising the steps of:

removing aluminate ions from the Bayer liquor through the formation of a carbonate-bearing hydrocalumite and/or sulphate-bearing hydrocalumite; and,

treating the liquor with sufficient lime to remove and causticise any residual carbonate ions and some or all of the oxalate ions present whereby any reacted lime solids thus formed can be separated and safely disposed of.

2. A process for the removal and causticisation of sodium oxalate and/or sodium sulphate as defined in claim 1, wherein the process comprises a further step, prior to said step of removing aluminate ions, in which the liquor is enriched with sulphate and/or oxalate such that any aluminate and/or carbonate ions entering with the sulphate and/or oxalate are also removed.

3. A process for the removal and causticisation of sodium oxalate and/or sodium sulphate as defined in claim 1, wherein the process comprises a further step, following said step of removing aluminate ions, of separating the carbonate-bearing hydrocalumite species and/or sulphate-bearing hydrocalumite species from the Bayer liquor to form a clarified liquor.

4. A process for the removal and causticisation of sodium oxalate and/or sodium sulphate as defined in claim 3, wherein the process comprises the further step of re-using said hydrocalumite species in an external causticisation process, whereby the aluminate contained within the hydrocalumite species is recovered.

5. A process for the removal and causticisation of sodium oxalate and/or sodium sulphate as defined in claim 1, wherein the process further comprises a pre-

causticisation step in which the Bayer liquor is first causticised to reduce the concentration of carbonate ions, prior to said step of removing aluminate ions.

6. A process for the removal and causticisation of sodium oxalate and/or sodium sulphate as defined in claim 5, wherein said pre-causticisation step includes heating the liquor to close to boiling point, adding sufficient lime to react with the carbonate ions to produce substantially calcium carbonate and separating the reacted lime solids from the liquor.
7. A process for the removal and causticisation of sodium oxalate and/or sodium sulphate as defined in claim 6, wherein the heated liquor is enriched with sulphate and/or oxalate prior to pre-causticisation to ensure that any carbonate ions entering with the sulphate and/or oxalate are also causticised.
8. A process for the removal and causticisation of sodium oxalate and/or sodium sulphate as defined in claim 1, wherein the S concentration of the Bayer process liquor prior to the said step of removing aluminate ions is between 0 and 250g/l, and the liquor is maintained at a temperature of between 20°C and 90°C, with a reaction time of up to 120 minutes.
9. A process for the removal and causticisation of sodium oxalate and/or sodium sulphate as defined in claim 8, wherein the S concentration of the Bayer process liquor prior to the said step of removing aluminate ions is less than 150 g/l, and the liquor is maintained at a temperature of between 50°C and 70°C, with a reaction time of approximately 30 minutes.
10. A process for the removal and causticisation of sodium oxalate and/or sodium sulphate as defined in claim 9, wherein said step of treating the clarified liquor with sufficient lime is conducted at a temperature between 20°C and 140°C, with a reaction time of between 0.25 to 4.0 hours.
11. A process for the removal and causticisation of sodium oxalate and/or sodium sulphate as defined in claim 10, wherein said step of treating the clarified liquor with

sufficient lime is conducted at a temperature between 50°C and 80°C, with a reaction time of between 0.5 to 1.0 hours.

12. A process for the removal and causticisation of sodium oxalate and/or sodium sulphate as defined in claim 11, wherein the said step of treating the liquor with sufficient lime is followed by a step of separating the reacted lime solids from the liquor and disposing of the solids or washing and drying the solids for calcination and re-use.

13. A process for the removal and causticisation of sodium oxalate and/or sodium sulphate as defined in claim 7, wherein the S concentration of the Bayer process liquor prior to said pre-causticisation step is between 0 and 250 g/l, and during said pre-causticisation step the liquor is heated to approximately 100°C.

14. A process for the removal and causticisation of sodium oxalate and/or sodium sulphate as defined in claim 13, wherein the clarified liquor is cooled to between 30°C and 90°C prior to said step of removing the aluminate ions.

15. An apparatus for the removal and causticisation of sodium oxalate and/or sodium sulphate from a Bayer process liquor containing sodium carbonate and one or both of sodium oxalate and sodium sulphate in an alumina refinery, the apparatus comprising:

means for removing aluminate ions from the Bayer liquor through the formation of a carbonate-bearing hydrocalumite and/or sulphate-bearing hydrocalumite; and,

means for treating the liquor with sufficient lime to remove and causticise any residual carbonate ions and some or all of the oxalate ions present whereby any reacted lime solids thus formed can be separated and safely disposed of.

16. An apparatus for removal and causticisation of sodium oxalate and/or sodium sulphate as defined in claim 15, wherein said means for removing aluminate ions comprises a first reaction vessel to which sufficient lime is added to react with all of

the aluminate ions in the liquor.

17. An apparatus for removal and causticisation of sodium oxalate and/or sodium sulphate as defined in claim 16, wherein the apparatus further comprises a means  
5 for separating the carbonate-bearing hydrocalumite and/or sulphate-bearing hydrocalumite from the liquor to form a clarified liquor.
18. An apparatus for removal and causticisation of sodium oxalate and/or sodium sulphate as defined in claim 17, wherein the means for treating the liquor comprises  
10 a second reaction vessel to which sufficient lime is added to react with the sodium oxalate in the clarified liquor to form calcium oxalate and with any remaining carbonate ions to form sodium carbonate.
19. An apparatus for removal and causticisation of sodium oxalate and/or sodium sulphate as defined in claim 15, wherein said means for removing aluminate ions  
15 and said means for treating the liquor are comprised in a single reaction vessel to which sufficient lime is added to react with the aluminate ions to form said carbonate-bearing and/or sulphate-bearing hydrocalumite, together with sufficient additional lime to react with the sodium oxalate to form calcium oxalate.  
20
20. An apparatus for removal and causticisation of sodium oxalate and/or sodium sulphate as defined in claim 19, wherein another first reaction vessel is provided for  
25 treating the liquor, prior to entry into said single reaction vessel, with sufficient lime to react with the sodium carbonate and then discharged to a solid/liquid separation device for separating any solids, which will consist primarily of calcium carbonate, to form a clarified liquor.

1/1

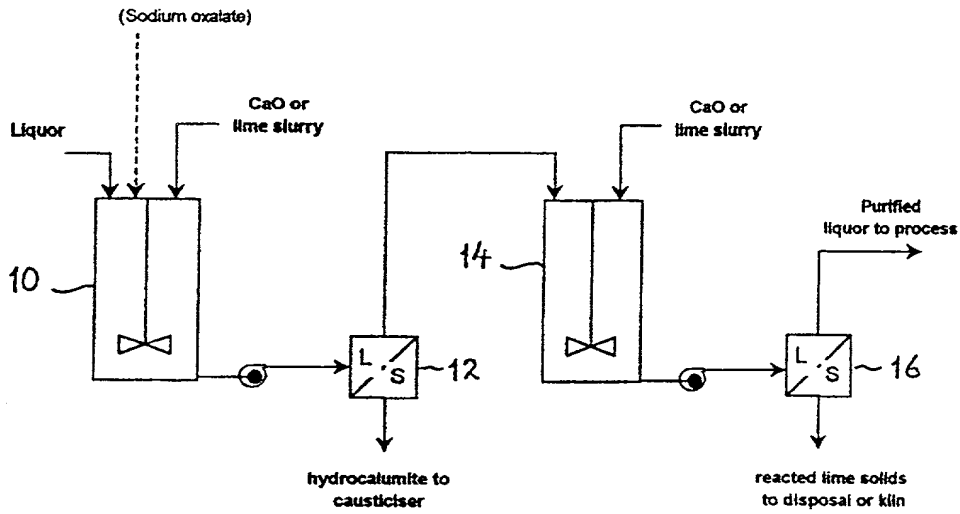


FIG. 1.

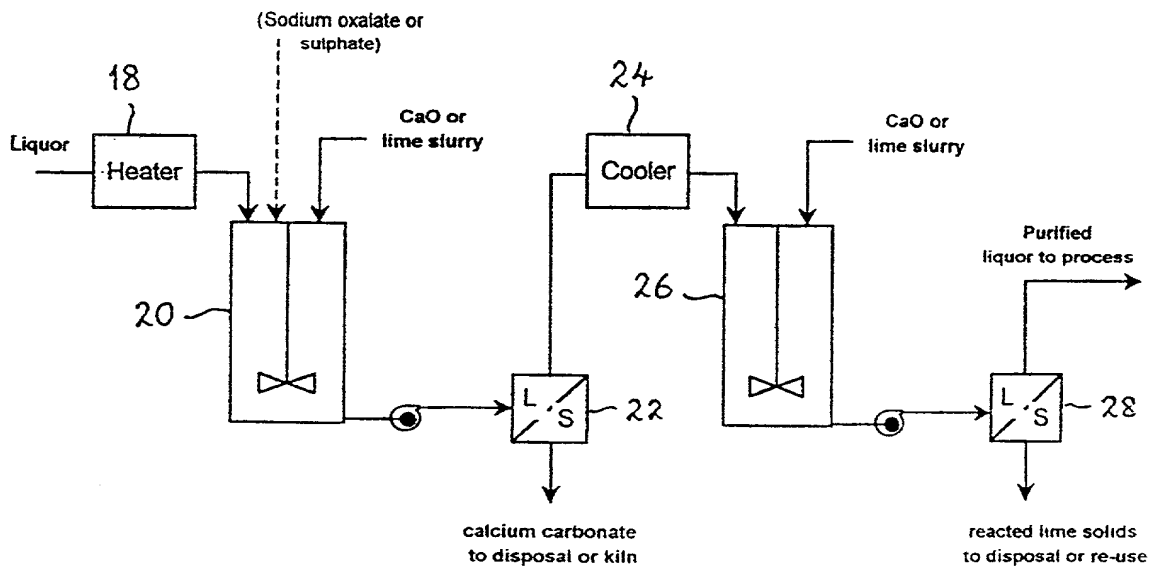


FIG. 2.

Please type a plus sign (+) inside this box → ☐

PTO/SB/01 (12-97)

Approved for use through 9/30/00. OMB 0651-0032

Patent and Trademark Office; U.S. DEPARTMENT OF COMMERCE

Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it contains a valid OMB control number.

<b>DECLARATION FOR UTILITY OR DESIGN PATENT APPLICATION (37 CFR 1.63)</b>	<b>Attorney Docket Number</b>	0730.0040C	
	<b>First Named Inventor</b>	Steven Rosenberg, et al	
	<b>COMPLETE IF KNOWN</b>		
	<b>Application Number</b>	/	
	<b>Filing Date</b>	Herewith	
	<b>Group Art Unit</b>	Not Yet Assigned	
<input checked="" type="checkbox"/> Declaration Submitted with Initial Filing	<input type="checkbox"/> Declaration Submitted after Initial Filing (surcharge (37 CFR 1.16 (e)) required)	<b>Examiner Name</b>	Not Yet Assigned

As a below named inventor, I hereby declare that:

My residence, post office address, and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

**Process for the Removal of Oxalate and/or Sulphate from Bayer Liquors**

the specification of which *(Title of the Invention)*

☒ is attached hereto  
OR  
☒ was filed on (MM/DD/YYYY) **March 17, 2000** as United States Application Number or PCT International Application Number **PCT/AU00/00208** and was amended on (MM/DD/YYYY) (if applicable).

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment specifically referred to above.

I acknowledge the duty to disclose information which is material to patentability as defined in 37 CFR 1.56.

I hereby claim foreign priority benefits under 35 U.S.C. 119(a)-(d) or 365(b) of any foreign application(s) for patent or inventor's certificate, or 365(a) of any PCT international application which designated at least one country other than the United States of America, listed below and have also identified below, by checking the box, any foreign application for patent or inventor's certificate, or of any PCT international application having a filing date before that of the application on which priority is claimed.

Prior Foreign Application Number(s)	Country	Foreign Filing Date (MM/DD/YYYY)	Priority Not Claimed	Certified Copy Attached?	
				YES	NO
PP 9334	Australia	03/19/1999	<input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/>	<input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/>	<input checked="" type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/>

☐ Additional foreign application numbers are listed on a supplemental priority data sheet PTO/SB/02B attached hereto.

I hereby claim the benefit under 35 U.S.C. 119(e) of any United States provisional application(s) listed below.

Application Number(s)	Filing Date (MM/DD/YYYY)

☐ Additional provisional application numbers are listed on a supplemental priority data sheet PTO/SB/02B attached hereto.

(Page 1 of 2)

Burden Hour Statement: This form is estimated to take 0.4 hours to complete. Time will vary depending upon the needs of the individual case. Any comments on the amount of time you are required to complete this form should be sent to the Chief Information Officer, Patent and Trademark Office, Washington, DC 20231. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Assistant Commissioner for Patents, Washington, DC 20231.

Please type a plus sign (+) inside this box → ☐PTO/SB/01 (12-97)  
Approved for use through 9/30/00. OMB 0651-0032  
Patent and Trademark Office, U.S. DEPARTMENT OF COMMERCE  
Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it contains a valid OMB control number.**DECLARATION — Utility or Design Patent Application**

I hereby claim the benefit under 35 U.S.C. 120 of any United States application(s), or 365(c) of any PCT international application designating the United States of America, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT international application in the manner provided by the first paragraph of 35 U.S.C. 112, I acknowledge the duty to disclose information which is material to patentability as defined in 37 CFR 1.56 which became available between the filing date of the prior application and the national or PCT international filing date of this application.

U.S. Parent Application or PCT Parent Number	Parent Filing Date (MM/DD/YYYY)	Parent Patent Number (if applicable)

☐ Additional U.S. or PCT international application numbers are listed on a supplemental priority data sheet PTO/SB/02B attached hereto.

As a named inventor, I hereby appoint the following registered practitioner(s) to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith:

☒ Customer Number 27896 OR ☐ Registered practitioner(s) name/registration number listed below

Name	Registration Number	Name	Registration Number

☐ Additional registered practitioner(s) named on supplemental Registered Practitioner Information sheet PTO/SB/02C attached hereto.

Direct all correspondence to: ☒ Customer Number 27896 OR ☐ Correspondence address below

Name			
Address			
Address			
City	State	ZIP	
Country	Telephone	Fax	

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under 18 U.S.C. 1001 and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Name of Sole or First Inventor: ☐ A petition has been filed for this unsigned inventor

Given Name (first and middle (if any))		Family Name or Surname	
Steven P.		Rosenberg	
Inventor's Signature	Date		13/9/01
Residence: City	Australind	State	WA
		Country	Australia
Post Office Address	16 Wallaroo Way		
Post Office Address			
City	Australind	State	WA
		ZIP	6233
		Country	Australia

☐ Additional inventors are being named on the supplemental Additional Inventor(s) sheet(s) PTO/SB/02A attached hereto

Please type a plus sign (+) inside this box →

PTO/SB/02A (3-97)

Approved for use through 9/30/98. OMB 0651-0032  
Patent and Trademark Office: U.S. DEPARTMENT OF COMMERCE

Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it contains a valid OMB control number.

DECLARATION				ADDITIONAL INVENTOR(S) Supplemental Sheet Page <u>1</u> of <u>1</u>				
Name of Additional Joint Inventor, if any:				<input type="checkbox"/> A petition has been filed for this unsigned inventor				
Given Name (first and middle [if any])				Family Name or Surname				
<u>Wayne</u>				<u>Tichbon</u>				
Inventor's Signature		<u>Wayne Tichbon</u>			Date		<u>14/9/2001</u>	
Residence: City		<u>Eaton</u>		State		<u>WA</u>		
				Country		<u>Australia</u> <u>AUX</u>		
				Citizenship		<u>AU</u>		
Post Office Address <u>9 Pinto Close</u>								
Post Office Address								
City		<u>Eaton</u>		State		<u>WA</u>		
				ZIP		<u>6232</u>		
				Country		<u>Australia</u>		
Name of Additional Joint Inventor, if any:				<input type="checkbox"/> A petition has been filed for this unsigned inventor				
Given Name (first and middle [if any])				Family Name or Surname				
<u>Darrel J.</u>				<u>Wilson</u>				
Inventor's Signature		<u>[Signature]</u>			Date		<u>14/9/01</u>	
Residence: City		<u>Leschenault</u>		State		<u>WA</u>		
				Country		<u>Australia</u> <u>AUX</u>		
				Citizenship		<u>AU</u>		
Post Office Address <u>33 Coral Drive</u>								
Post Office Address								
City		<u>Leschenault</u>		State		<u>WA</u>		
				ZIP		<u>6233</u>		
				Country		<u>AU</u>		
Name of Additional Joint Inventor, if any:				<input type="checkbox"/> A petition has been filed for this unsigned inventor				
Given Name (first and middle [if any])				Family Name or Surname				
<u>Catherine A.</u>				<u>Heath</u>				
Inventor's Signature		<u>C.A. Heath</u>			Date		<u>14/9/01</u>	
Residence: City		<u>East Bunbury</u>		State		<u>WA</u>		
				Country		<u>Australia</u> <u>AUX</u>		
				Citizenship		<u>AU</u>		
Post Office Address <u>36A Moore Street</u>								
Post Office Address								
City		<u>East Bunbury</u>		State		<u>WA</u>		
				ZIP		<u>6230</u>		
				Country		<u>Australia</u>		

**Burden Hour Statement:** This form is estimated to take 0.4 hours to complete. Time will vary depending upon the needs of the individual case. Any comments on the amount of time you are required to complete this form should be sent to the Chief Information Officer, Patent and Trademark Office, Washington, DC 20231. **DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO:** Assistant Commissioner for Patents, Washington, DC 20231.